SUBSTITUTE FORM PTO-1449 U.S. DEPARTMENT OF COMMERCE	ATTY. DOCKET NO. JEFF-YOO01	SERIAL NO. 09/473,872
	APPLICANT: Yoon	
	FILING DATE 12/28/99	GROUP 1632

U.S. PATENT DOCUMENTS

EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE (IF APPROPRIATE)

FOREIGN PATENT DOCUMENTS

		DOCUMENT NO.	DATE OF PUBLICATION	COUNTRY	CLASS	SUBCLASS	TRANSLATION YES	NO

OTHER DOCUMENTS ***(Including Author, Title, Date, Pertinent Pages, Etc.)***

9w	AA	√	Ahmad et al., "Alopecia Universalis Associated with a Mutation in the Human *hairless* Gene," Science, 1998, Vol. 279, p. 720-724
	AB	√	Armstrong et al., "Haploinsufficiency of desmoplakin causes a striate subtype of palmoplantar keratoderma", Human Molecular Genetics," 1999, Vol. 8, No.1, p. 143-148
	AC	√	Dong et al., "Frequent Somatic Mutations in *Serine/Threonine Kinase 11*/Peutz-Jeghers Syndrome Gene in Left-sided Colon Cancer," Cancer Research, 1998, Vol. 58, p. 3787-3790
	AD	√	Frank et al., "Exposing the human *nude* phenotype," Nature, 1999, Vol. 398, p. 473-474
	AE	√	Korge et al., "A Mutational Hotspot in the 2B Domain of Human Hair Basic Keratin 6 (hHb6) in Monilethrix Patients," 1998, Vol. 111, No. 5, p 896-898
	AF	√	Marsh et al., "Germline mutations in *PTEN* are present in Bannayan-Zonana syndrome," Nature Genetics, 1997, Vol. 16, p. 333-334
	AG	√	McGrath et al., "Mutations in the plakophilin 1 gene result in ectodermal dysplasia/skin fragility syndrome,", Nature Genetics, 1997, Vol. 17, p. 240-244
	AH	√	Richard et al., "Functional defects of Cx26 resulting from a heterozygous missense mutation in a family with dominant deaf-mutism and palmoplantar keratoderma," Hum Genet, 1998, Vol. 103, p. 393-399
	AI	√	Richard et al., "Mutations in the human connexin gene *GJB3* cause erythrokeratodermia variabilis," Nature Genetics, 1998, Vol. 20, p. 366-369
	AJ	√	Rickman et al., "N-terminal deletion in a desmosomal cadherin causes the autosomal dominant skin disease striate palmoplantar keratoderma," Human Molecular Genetics, 1999, Vol. 8, No. 6, p. 971-976
	AK	√	Rowan et al., "Somatic Mutations in the Peutz-Jegners (LKB1/STKII) Gene in Sporadic Malignant Melanomas," The Journal of Investigative Dermatology, 1999, Vol. 112, No. 4, p. 509-511
	AL	√	Sakuntabhai et al., "Mutations in *ATP2A2*, encoding a Ca^{2+} pump, cause Darier disease," Nature Genetics, 1999, Vol. 21, p. 271-277
	AM	√	Winter et al., "Mutations in the hair cortex keratin hHb6 cause the inherited hair disease monilethrix," Nature Genetics, 1997, Vol. 16, p. 372-374

EXAMINER Joe Woitach	DATE CONSIDERED 11/28/01

EXAMINER: Initial if citation considered, whether or not citation is in conformance with MPEP 609; Draw line through Citation if not in conformance and not considered. Include copy of this form with next communication to applicant.